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- A hybridoma selected from 5F8 (ECACC No.95121524), 2H6 (ECACC No.95121526) and 5D8 (ECACC No.95121527).
- 16. A monoclonal antibody selected from the monoclonal antibodies produced by the hybridomas according to claim 15.



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	NPL	CTNF
APPL PARTS	Non-Patent Literature	Count Non-Final
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nternal Misc. Paper	PET	EXIN
LET:	Petition Petition	Examiner Interview
Aisc. Incoming Letter	RETMAIL	M903
371P	Mail Returned by USPS	DO/EO Acceptance
•	SEQLIST	M905
Amendment Including Elections	Sequence Listing	M905 DO/EO Missing Requirement
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ABST	Specification SPEC	Formal Drawing Required
Abstract		NOA
ADS	SPEC NO	Notice of Allowance
Application Data Sheet	TRNA	PETDEC
AF/D Affidavit or Exhibit Received	Transmittal New Application	Petition Decision
Appendix Appendix		
, ,	OUTCOING	INCOMING
Artifact ARTIFACT	OUTGOING	MCOMING
	CTMS	AP.B
Bib Data Sheet	Misc. Office Action	Appeal Brief
_ CLM	1449	C.AD Change of Address
Claim CLIVI	Signed 1449	
COMPUTER	892	N/AP
Computer Program Listing	892	Notice of Appeal
CRFL	ABN	PA Change in Power of Attorney
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All CRF Papers for Backfile	APDEC	REM
DIST Terminal Disclaimer Filed	Board of Appeals Decision	Applicant Remarks in Amendmen
	APEA	XT/
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SRNT	WCLM	IIFW
Examinor Search Notes	Claim Worksheet	File Wrapper Issue Information

WFEE

SRFW File Wrapper Search Info

Fee Worksheet

Examiner Search Notes

CLMPTO _______ PTO Prepared Complete Claim Set

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(71)(72) Applicant and Inventor: MÅRDH, Sven Sjöliden 13, S-590 77 Vreta Kloster (SE).	[SE/SI	TO TO THE STATE OF	
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(SASTIMAL DECOMBINANT PHACES			

(54) Title: RECOMBINANT PHAGES

(57) Abstract

The present invention relates to bacteriophages for use in the treatment or prophylaxis of bacterial infections, especially mucosal bacterial infections such as *Helicobacter pylori* infections. In particular, it relates to modified filamentous bacteriophages, e.g. M13 phages, for such use, which bacteriophages present at its surface a recombinant protein comprising: (i) a first component derived from a bacteriophage surface protein; and (ii) a second component comprising variable region sequences of an antibody to provide a bacterial antigen binding site, said second component rendering said bacteriophage capable of binding to and thereby inhibiting growth of bacterial cells involved in the etiology of said infection.